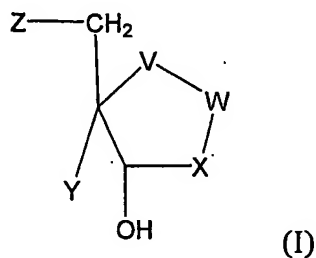


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Amendments to the Claims:

Please cancel Claim 10, 19, 26 and 28 without prejudice or disclaimer, amend Claims 1, 11, 23-25 and 27, and add new Claims 29-31 as set forth below.

1. (Currently amended) A compound of the formula (I):



wherein:

V is selected from CH<sub>2</sub> and NH, and W is selected from NR<sup>1</sup> and NR<sup>2</sup>; or V is selected from NR<sup>1</sup> and NR<sup>2</sup>, and W is selected from CH<sub>2</sub> and NH;

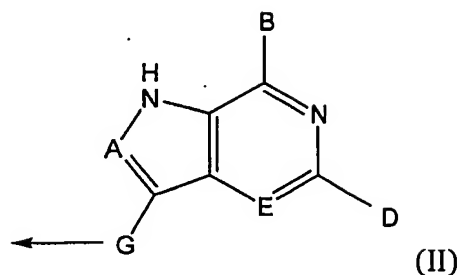
X is selected from CH<sub>2</sub> and CHOH in the R or S-configuration;

Y is selected from hydrogen, halogen and hydroxy, except where V is selected from NH[[,]] and NR<sup>1</sup> ~~and~~ NR<sup>2</sup> then Y is hydrogen;

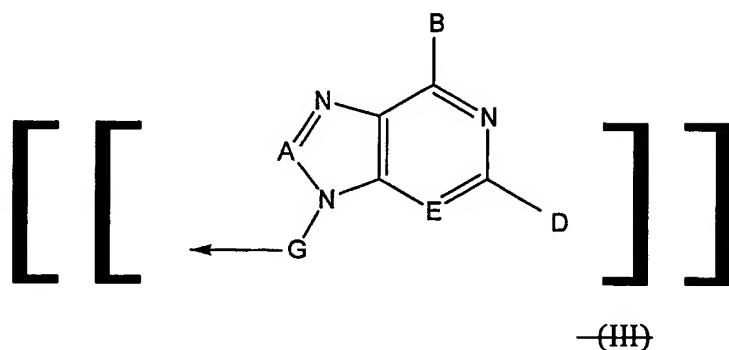
Z is selected from hydrogen, halogen, hydroxy, SQ, OQ and Q, where Q is an optionally substituted alkyl, aralkyl or aryl group;

R<sup>1</sup> is a radical of the formula (II)

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~~R<sup>3</sup> is a radical of the formula (III)~~



A is selected from N, CH and CR, where R is selected from halogen, optionally substituted alkyl, aralkyl or aryl, OH, NH<sub>2</sub>, NHR<sup>3</sup>, NR<sup>3</sup>R<sup>4</sup> and SR<sup>5</sup>, where R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> are each optionally substituted alkyl, aralkyl or aryl groups;

B is selected from OH, NH<sub>2</sub>, NHR<sup>6</sup>, SH, hydrogen and halogen, where R<sup>6</sup> is an optionally substituted alkyl, aralkyl or aryl group;

D is selected from OH, NH<sub>2</sub>, NHR<sup>7</sup>, hydrogen, halogen and SCH<sub>3</sub>, where R<sup>7</sup> is an optionally substituted alkyl, aralkyl or aryl group;

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E is selected from N and CH;

G is selected from CH<sub>2</sub> and NH, or G is absent, provided that where W is NR<sup>1</sup> ~~or NR<sup>2</sup>~~ and G is NH then V is CH<sub>2</sub>, and provided that where V is NR<sup>1</sup> ~~or NR<sup>2</sup>~~ and G is NH then W is CH<sub>2</sub>,

or a tautomer thereof, or a pharmaceutically acceptable salt thereof, ~~or an ester thereof,~~  
~~or a prodrug thereof.~~

2. (Previously presented) A compound as claimed in claim 1, where Z is selected from hydrogen, halogen, hydroxy, SQ and OQ.
3. (Previously presented) A compound as claimed in claim 1, where V is CH<sub>2</sub>.
4. (Previously presented) A compound as claimed in claim 1, where X is CH<sub>2</sub>.
5. (Previously presented) A compound as claimed in claim 1, where G is CH<sub>2</sub>.
6. (Previously presented) A compound as claimed in claim 1, where Z is OH.
7. (Previously presented) A compound as claimed in claim 1, where Z is SQ.
8. (Previously presented) A compound as claimed in claim 1, where Z is Q.
9. (Previously presented) A compound as claimed in claim 1, where W is NR<sup>1</sup>.

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10. (Canceled)

11. (Currently amended) A compound as claimed in claim 1, where W is selected from NH and NH,  $\text{NR}^1$  ~~or~~  $\text{NR}^2$  and X is  $\text{CH}_2$ .

12. (Previously presented) A compound as claimed in claim 1, where V, X and G are all  $\text{CH}_2$ , Z is OH and W is  $\text{NR}^1$ .

13. (Previously presented) A compound as claimed in claim 1, where V, X and G are all  $\text{CH}_2$ , Z is SQ and W is  $\text{NR}^1$ .

14. (Previously presented) A compound as claimed in claim 1, where Y is hydrogen.

15. (Previously presented) A compound as claimed in claim 1, where Y is hydroxy.

16. (Previously presented) A compound as claimed in claim 1, where B is hydroxy.

17. (Previously presented) A compound as claimed in claim 1, where B is  $\text{NH}_2$ .

18. (Previously presented) A compound as claimed in claim 1, where A is CH.

19. (Canceled)

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20. (Previously presented) A compound as claimed in claim 1, where D is H.

21. (Previously presented) A compound as claimed in claim 1, where D is NH<sub>2</sub>.

22. (Previously presented) A compound as claimed in claim 1, where E is N.

23. (Currently amended) A compound as claimed in claim 1, which is:

(3R,4R)-1-[(9-deazahypoxanthin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)pyrrolidine;

(3R,4R)-1-[(9-Deazaadenin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)pyrrolidine;

~~(3R,4R)-1-[(8-aza-9-deazahypoxanthin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)~~  
pyrrolidine;

~~(3R,4R)-1-[(8-aza-9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)pyrrolidine;~~

(3R,4S)-1-[(9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(2-phenylethyl)pyrrolidine;

(3S,4R)-1-[(9-deazahypoxanthin-9-yl)methyl]-3,4-dihydroxy-4-methylthiomethyl  
pyrrolidine;

(3R,4S)-1-[(9-deazahypoxanthin-9-yl)methyl]-3-hydroxy-4-(methylthiomethyl)  
pyrrolidine;

N-(9-Deazahypoxanthin-9-yl)-1,4-dideoxy-1,4-imino-D-ribitol;

N-(9-deazahypoxanthin-9-yl)methyl-1,4-dideoxy-1,4-imino-D-ribitol;

~~(3R,4R)-3-hydroxy-4-hydroxymethyl-1-(hypoxanthin-9-yl)pyrrolidine;~~

(3R,4S)-1-[(9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(methylthiomethyl)pyrrolidine;

(3R,4S)-1-[(9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(benzylthiomethyl)pyrrolidine;

~~(3R,4S)-1-[(8-aza-9-deezaadenin-9-yl)methyl]-3-hydroxy-4-(benzylthiomethyl)~~  
pyrrolidine;

(3R,4R)-1-[(9-deazaguanin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)pyrrolidine;

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(3R,4S)-1-[(9-deazaadenin-9-yl)methyl]-3-hydroxy-4-(4-chlorophenylthiomethyl)  
pyrrolidine;

(3R,4R)-1-[(6-chloro-9-deazapurin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)  
pyrrolidine;

(3R,4R)-1-[(6-azido-9-deazapurin-9-yl)methyl]-3-hydroxy-4-(hydroxymethyl)  
pyrrolidine; or

(3R,4R)-1-[(9-deazaadenin-9-yl)methyl]-3-acetoxy-4-(acetoxymethyl)pyrrolidine;  
or a pharmaceutically acceptable salt thereof, ~~or an ester thereof, or a prodrug thereof.~~

24. (Currently amended) A pharmaceutical composition comprising a  
pharmaceutically effective amount of a compound as claimed in claim 1 and a carrier.

25. (Currently amended) A method of treating a subject having a disease or  
condition in which it is desirable to inhibit purine phosphoribosyltransferase, purine  
nucleoside phosphorylase, 5'-methylthio adenosine phosphorylase, 5'-  
methylthioadenosine nucleosidase and/or nucleoside hydrolase comprising  
administering a ~~pharmaceutically effective amount of a~~ compound as claimed in claim 1  
to the subject in an amount effective to inhibit purine phosphoribosyltransferase, purine  
nucleoside phosphorylase, 5'-methylthio adenosine phosphorylase, 5'-  
methylthioadenosine nucleosidase and/or nucleoside hydrolase, a patient requiring  
~~treatment. wherein the disease or condition is a cancer, a bacterial infection, a protozoal~~  
infection, a T-cell mediated disease or a transplant rejection.

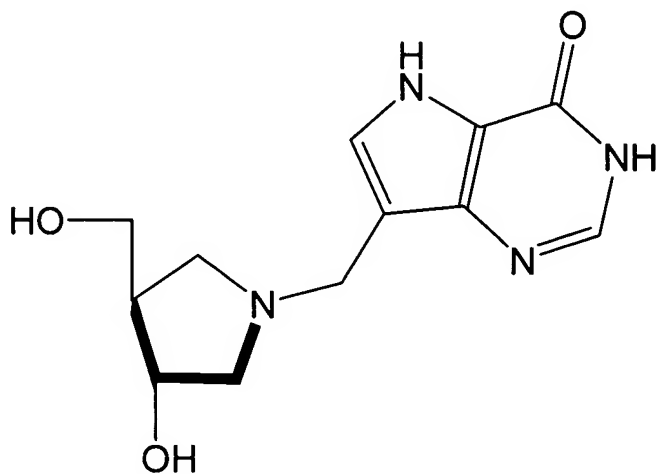
26. (Canceled)

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27. (Currently amended) The method of claim 25, ~~26~~, where the T-cell mediated disease is psoriasis or [[,]] arthritis ~~or transplant rejection~~.

28. (Canceled)

29. (New) The compound of claim 1 having the structure:



or a pharmaceutically acceptable salt thereof.

30. (New) The compound of claim 1, wherein the salt is derived from an acid selected from the group consisting of hydrochloric acid, sulphuric acid, phosphoric acid, acetic acid, lactic acid, fumaric acid, succinic acid, tartaric acid, gluconic acid, citric acid, methanesulfonic acid and p-toluenesulfonic acid.

31. (New) The compound of claim 29, wherein the salt is a hydrochloride salt.